

ISRE (Luc) HEK293 Reporter Cell Data Sheet

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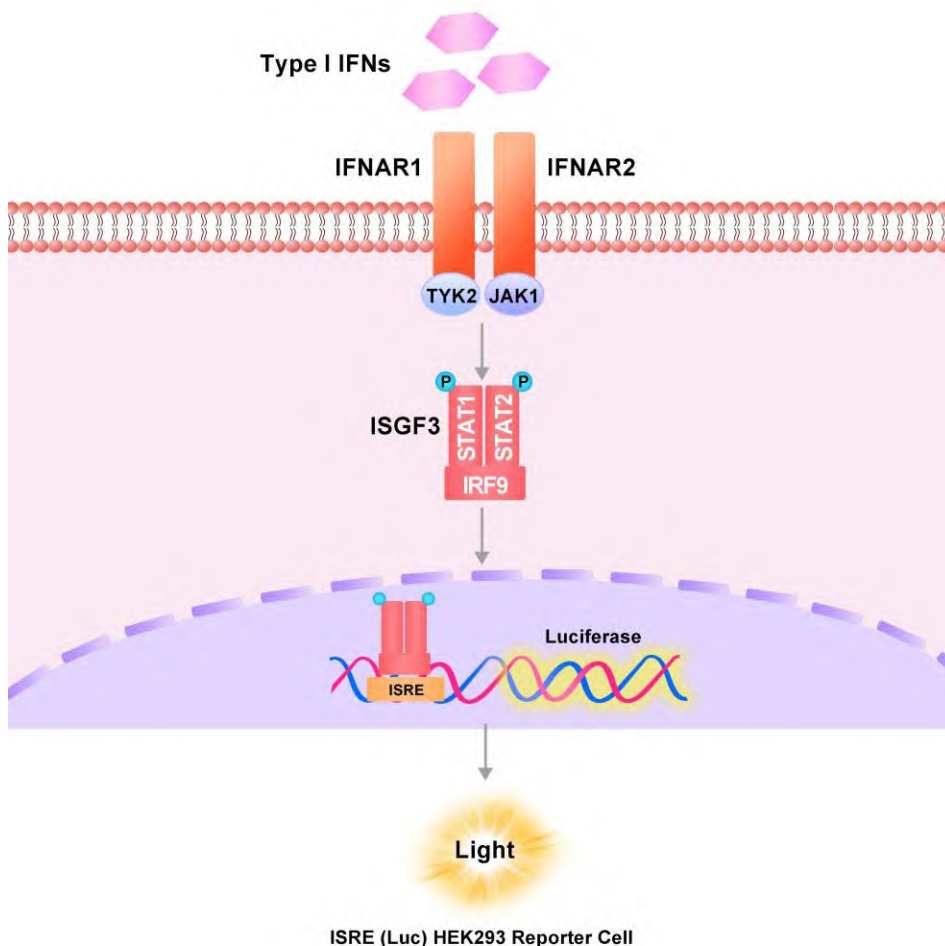
Catalog No.	Size
CHEK-ATF134	2 × (1 vial contains ~5×10 ⁶ cells)

• *Description*

The ISRE (Luc) HEK293 Reporter Cell was engineered to express IFN-stimulated response elements (ISRE) driving luciferase expressing systems. When stimulated with human Type I interferons (IFNs), receptor-mediated signaling can drive IFN-stimulated gene factor 3 (ISGF3)-mediated luminescence.

• *Application*

- Screen for activators or inhibitors of Type I interferons.



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• Cell Line Profile

Cell line	ISRE (Luc) HEK293 Reporter Cell
Host Cell	HEK293
Property	Adherent
Complete Growth Medium	DMEM + 10% FBS
Selection Marker	Puromycin (2 µg/mL)
Incubation	37°C with 5% CO ₂
Doubling Time	22-24 hours
Transduction Technique	Lentivirus

• Materials Required for Cell Culture

- DMEM medium (Gibco, Cat.No.11965-092)
- Fetal bovine serum (CellMax, Cat.No.SA211.02)
- Puromycin (InvivoGen, Cat.No.ant-pr-5b)
- Complete Growth Medium: DMEM + 10% FBS
- Culture Medium: DMEM + 10% FBS, Puromycin (2 µg/mL)
- Freeze Medium: 90% FBS, 10% (V/V) DMSO
- T-75 Culture flask (Corning, 430641)
- Cryogenic storage vials (SARSTEDT, 72.379.007)
- Thermostat water bath
- Centrifuge
- Luna cell counter (Logos Biosystems, LUNA- II)
- CO₂ Incubator (Thermo, 3111)
- Biological Safety Cabinet (Thermo, 1389)

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• *Recovery*

1. Thaw the vial by gentle agitation in a 37°C water bath. To reduce the possibility of contamination, keep the cap out of the water. Thawing should be rapid (approximately 2 minutes).
2. Remove the vial from the water bath as soon as the contents are thawed, and decontaminate by spraying with 70% ethanol. All the operations from this point on should be carried out under strict aseptic conditions.
3. Transfer the vial contents to a centrifuge tube containing 4.0 mL complete growth medium and spin at approximately 1000 rpm for 5 minutes.
4. Discard the supernatant and resuspend cell pellet with 5 mL complete growth medium and transfer the cell suspension into T-75 flask containing 10-15 mL of pre-warmed complete growth medium.
5. Incubate at 37°C with 5% CO₂ incubator.

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• *Subculture*

1. Remove and discard culture medium.
2. Wash the cells once with sterile PBS.
3. Add 2 mL of 0.25% trypsin to cell culture flask. Place the flask at 37°C for 2-3 minutes, until 90% of the cells have detached.
4. Add 6.0 to 8.0 mL of culture medium and aspirate cells by gently pipetting.
5. Add appropriate aliquots of the cell suspension to new culture vessel.
6. Incubate at 37°C with 5% CO₂ incubator.
 - **Subcultivation Ratio:** A subcultivation ratio of 1:6 to 1:10 is recommended.
 - **Medium Renewal:** Every 2 to 3 days.

• *Cryopreservation*

1. Remove and discard spent medium.
2. Detach cells from the cell culture flasks with 0.25% trypsin.
3. Centrifuge at 1000 rpm for 5 min at RT to pellet cells.
4. Resuspend the cell pellets with complete growth medium and count viable cells.
5. Centrifuge at 1000 rpm for 5 min at RT and resuspend cells in freezing medium to a concentration of 5×10^6 to 1×10^7 cells/mL.
6. Aliquot into cryogenic storage vials. Place vials in a programmable cooler or an insulated box placed in a -80°C freezer overnight, then transferring to liquid nitrogen storage.

• *Storage*

- **Product format:** Frozen
- **Storage conditions:** Liquid nitrogen immediately upon receipt

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• *Signaling Bioassay*

Human IFN-alpha 2b Protein Stimulation (RLU)

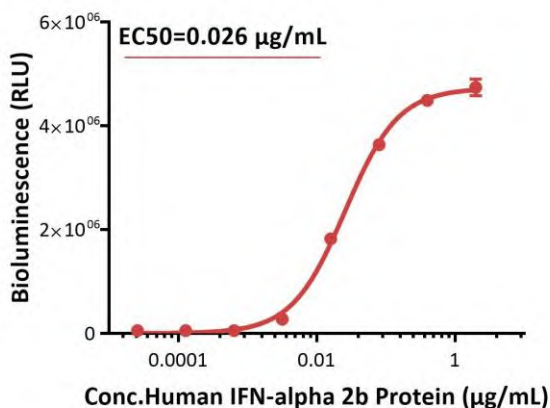


Fig1. Response to human IFN-alpha 2b protein (RLU). The ISRE (Luc) HEK293 Reporter Cell was stimulated with serial dilutions of human IFN-alpha 2b protein (Cat.No.IFB-H5253). The EC50 was approximately 0.026 $\mu\text{g/mL}$.

Human IFN-alpha 2b Protein Stimulation (FOLD)

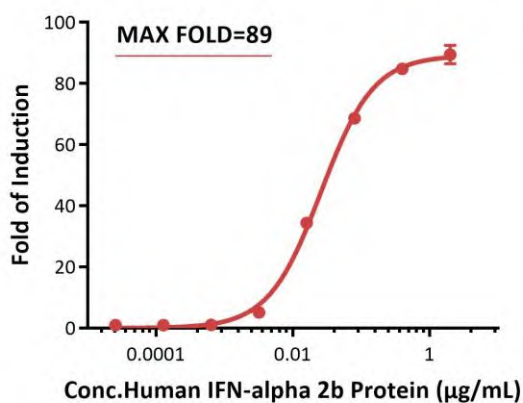


Fig2. Response to human IFN-alpha 2b protein (FOLD). The ISRE (Luc) HEK293 Reporter Cell was stimulated with serial dilutions of human IFN-alpha 2b protein (Cat.No.IFB-H5253). The max induction fold was approximately 89.

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• *Related Products*

Products

Human IFN-alpha 2b Protein

Cat.No.

IFB-H5253